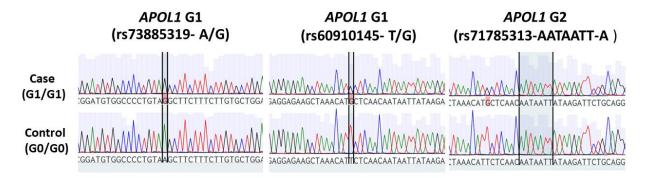
Supplementary material:

Supplementary methods: APOL1 Genotyping

DNA was extracted from paraffin embedded tissue from kidney biopsy using the protocol described in QIAamp DNA FFPE Tissue Kit Manual. The *APOL1* G1 risk allele is represented by both rs73885319 and rs60910145 missense variants, and the G2 risk allele is a 6 bp deletion (rs71785313, formerly rs143830837)⁷. Genotyping for the APOL1 G1 and G2 variants was conducted using PCR amplification and Sanger sequencing. Analysis of sequence data demonstrated that the patient was homozygous for the G1 allele.

Supplementary figure:

Chromatogram of rs73885319 and rs60910145 sequence, representing the linked missense variants for the *APOL1* G1 risk allele, and rs71785313 sequence, representing the 6 bp deletion for the *APOL1* G2 risk allele. The patient is homozygous for the G1 allele.



Supplementary References:

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